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(54) APPARATUS FOR PROMOTING SOFT TISSUE ENLARGEMENT AND WOUND HEALING

**VORRICHTUNG ZUR FÖRDERUNG DER WEICHGEWEBEVERGRÖßERUNG UND DER
WUNDHEILUNG**

**APPAREIL DESTINE A FAVORISER L'ELARGISSEMENT DE TISSUS MOUS ET LA
CICATRISATION DES PLAIES**

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EP 0 708 620 B1

Description

Cross Reference to Related Application

[0001] This application is a continuation-in-part of Serial No: 08/220,186 filed March 30, 1994, now U.S. Pat. No. 5,701,917, issued December 30, 1997.

Background of the Invention

[0002] There are numerous instances where persons desire enlargement of the soft tissues in their bodies. One such instance is for the replacement of one or both breasts amputated during a mastectomy in order to restore physiological symmetry and psychological well-being. Other instances are for correction of natural abnormalities such as dimpling. Still other instances are for augmentation of physical attributes to improve cosmetics and self-esteem. These latter soft tissue enlargements are principally directed to breast enlargement in females and penis enlargement in males.

[0003] Prosthetic implants have been developed for insertion below the skin. However, the severity of the potential complications including scarring, implant rupture, capsular contracture, necrosis and implant migration as well as the recite adverse publicity thereof have significantly reduced the desirability of these implants. Thus, there is a societal need for other means to obtain soft tissue enlargement.

[0004] Some soft tissue enlargements occur naturally. For instance, during pregnancy the skin over a woman's abdominal region enlarges approximately nine times its previous area to accommodate the fetus without a proportional decrease in skin thickness. In other words, the abdominal skin tissue actually enlarges and does not merely stretch during pregnancy. Similarly, the skin will expand to accommodate any growth under the skin.

[0005] In the past, plastic surgeons have used this phenomena to their advantage to expand skin in order to accommodate prosthetic implants or provide tissue to close wound defects. To conduct this procedure, the surgeon inserts a balloon beneath the skin in the area where additional skin is desired. By progressively expanding the balloon, the skin first stretches and eventually actually grows to accommodate the increased volume underneath it. When the desired amount of skin is formed, the balloon is deflated and removed, and the implant is inserted into the cavity left by the balloon. Also, the excess skin can be used to cover a wound defect, an ulcer or a depressed scar. Similar methods have been used by African native tribes to enlarge lips, nostrils, and earlobes.

[0006] Other surgical techniques have used tissue expansion to achieve other types of soft tissue growth. For instance, balloons have been successfully situated underneath nerves, veins, tendons, and the like to expand and thereby elongate these tissues to repair damage

and alleviate various abnormalities.

[0007] A more advanced surgical method is known as callotasis distraction osteogenesis or limb lengthening. This method comprises cutting the bone about its periphery at the location where lengthening is desired, leaving the tissues inside and around the bone intact. Brackets are attached to the bone on each side of the separation, and the bone segments are slowly pulled away from one another while remaining integral over a period of several months. Not only does this cause the mended bone to be longer, but also the soft tissue surrounding the bone actually grows to accommodate the increased limb length. Similar methods have been used by African native tribes to lengthen necks for cosmetic purposes.

[0008] Each of these above-mentioned apparatuses and methods requires an invasive surgical technique to accomplish the soft tissue expansion. Invasive techniques increase the likelihood of the complications associated with the procedure including those mentioned above with respect to implant surgery. In addition, the expense of surgery precludes many persons having their abnormalities corrected or physical attributes enhanced.

[0009] Other soft tissue enlargement techniques have been developed which use other mechanisms to cause the enlargement. For instance, an instrument and technique have been developed for the non-surgical correction of inverted nipples due to short lactiferous ducts. The instrument is comprised of a cup having an internal volume shaped like that of the final desired nipple. The user places the cup over the inverted nipple, pumps the air out of the cup with a syringe and adjusts the vacuum within the cup using a check valve to just below the threshold of discomfort. Thus attached, the device puts the lactiferous ducts in tension and extends them sufficiently after two to three months of wear at 8-12 hours per day.

[0010] Although this device is sufficient for its intended purpose, it is not suitable for general soft tissue enlargement. Laceration and contusion can occur if too strong of a suction is applied to soft tissue. As the pressure within the inverted nipple instrument is not regulated, contusion or laceration can occur. When a vacuum is developed within the cup of the instrument, an equal and opposite force is applied to the patient about the rim of the cup. Excessive contact forces against the patient can cause ulceration, laceration, and contusions. As the contact forces are not regulated in the nipple instrument, these further complications also can occur. In addition, general soft tissue enlargement is not feasible with the instrument due to the size and shape of the cup.

[0011] Another prior art device is disclosed in U.S. Patent No. 936,434 as a device for enlarging a woman's breasts. This device included a pair of cups for placement on the breasts and a pump for exhausting the air. However, this patent provides no teaching as to the pressures to be used, the potential danger to the skin

tissues, or any suggestions as to how the device is to be retained in place during use. Apparently, the device is used in a clinical setting and is not suitable for long term ambulatory wear such as for 8-10 hours. As the patent suggests that the vacuum acts to cause the veins and arteries to engorge, thereby nourishing the breasts, it is clear that the patentee is suggesting that the breast tissue actually expands through this expansion of blood vessels alone. This patent has been the subject of ridicule by at least one medical authority. See "An Anthology Of Plastic Surgery" edited by Harry Hayes, Jr., M. D., Section 6, "Quackery and Nostrums" pub. 1986 by Aspen Publishers, Rockville, Maryland.

[0012] Finally, another prior art device although notorious is worthy of note. This device is commonly referred to as a penis pump and is sold primarily as a novelty as its long-term enlargement efficacy has never been proven and is in fact universally disclaimed by its distributors. The device is comprised of a cylinder having one open end into which the penis is inserted and a pump attached to it such that a vacuum can be created within the cylinder. Not only does this device have the same drawbacks as the nipple instrument with respect to potential complications, but also it is unlikely that sufficient vacuum can be maintained by the device to cause any notable long-term soft tissue enlargement. Further, this device is apparently designed to accomplish two tasks unrelated to enlargement. First, the device is used for stimulation and sexual gratification. Second, the device is used to promote erection by drawing blood into the penis.

[0013] There is also another condition routinely experienced by many patients in which the generation of soft tissue is important. That condition evolves from injuries or diseases which produce wound infections or ulcers which have a tendency to exude bodily fluids and resist healing. At least one effort has been made in the prior art, as known by the inventor herein, to address this problem. This prior art solution involves the use of an occlusive, or airtight, dressing covering the wound coupled with a suctioning of fluid from the wound either once or repeatedly in order to dry it out and create an environment more conducive to wound healing. It is not believed that this prior art considers the problems of excessive contact forces against the patient's skin which can cause ulceration, laceration, and contusions. Also, this prior art teaching is not focused on soft tissue enlargement through the use of a vacuum alone and instead relies at least in part on the use of suction for removing wound fluid and creating an environment that promotes healing. The importance of enlarging the surrounding soft tissue to close the wound is not a clear focus of this prior art method.

[0014] Most of these prior art devices and methods have failed to achieve long term soft tissue enlargement while preventing damage to the soft tissue being enlarged, as well as surrounding tissue. The inventor herein has succeeded in designing and developing a new

generalized method and apparatus for soft tissue enlargement which prevents damage to soft tissue. The apparatus used for this enlargement is comprised of a rigid fluid-impervious dome having a rim about its periphery and a vacuum pump for reducing pressure to thereby apply a distracting force to the soft tissue isolated by and within the dome. The rim has sufficient surface area such that the pressure applied to the patient by the rim is less than or equal to the negative pressure applied to the soft tissue under the dome. Thus, as long as pressure within the dome is regulated to a limit below which medical complications cannot occur, the opposing contact pressure against the patient is below this threshold as well. With this approach, damage is avoided not only to the soft tissue being enlarged, but the surrounding tissue as well. In the preferred embodiment of the apparatus, the vacuum pump has a self-contained power source. In addition, a pressure sensor and servomechanism control the pump such that the vacuum within the dome is maintained at a magnitude less than 35 mm Hg. Variant embodiments may be configured to fit over and enlarge a human breast, a human penis, an infected wound, open sore, ulcer, or any other desired area.

[0015] The method of use is comprised of the steps of attaching the dome to the location of desired enlargement, and creating a vacuum within the dome. The vacuum should be maintained for a minimum of eight hours per day and results should be sufficient after several months.

[0016] While the practical advantages and features of the present invention and method have been briefly described above, a greater understanding of the novel and unique features of the invention may be obtained by referring to the drawings and Detailed Description of the Preferred Embodiment which follow.

Brief Description of the Drawings

[0017]

Figure 1 is a front elevation view of the soft tissue enlargement apparatus of the present invention, showing the breast augmentation embodiment;

Figure 2 is a cross-sectional view of the breast enlargement embodiment taken in the plane of line 2-2 of Figure 1;

Figure 3 is a cross-sectional schematic of a dome and soft tissue in the early stages of enlargement;

Figure 4 is a cross-sectional schematic of a dome and soft tissue in the latter stages of enlargement;

Figure 5 is an orthographic projection of the penile augmentation embodiment of the present invention; and

Figure 6 is a partial cross-sectional view of a dome in place over an open wound.

Detailed Description of the Preferred Embodiment

[0018] The soft tissue enlargement apparatus 10 is generally comprised of a dome 12 having a rim 14 and vacuum pump assembly 16 for creating a vacuum within the dome. Although the vacuum pump assembly 16 may be a separate hand-held pump in one variant embodiment, in the preferred embodiment the vacuum pump assembly 16 is a self-contained vacuum pump 20 with an independent power source 22, pressure sensor 24, and servomechanism 26 for driving, regulating and controlling the vacuum pump 20.

[0019] Regulation of the pressure within the dome is essential to prevent contusions caused by rupturing capillaries adjacent the surface of the skin, separating epidermis from dermis and causing blisters. Medical data suggest that these contusions and blisters will not occur if pressure within the dome is maintained at less than 25-35 mm Hg for extended periods of time. Thus, the vacuum pump 20 must be regulated to control the pressure within the dome to within this limit. In addition, skin ulceration can occur if excessive contact pressures are applied thereto. Medical data suggest that a contact pressure less than 15-20 mm Hg may be applied indefinitely without such ulceration. However, contusions may occur due to positive contact pressures upon the skin at pressures for appreciable time periods above this ulceration limit. The preferred embodiment of the present invention was developed with these limits in mind and will not apply a continuous vacuum or a continuous contact pressure greater than 25-35 mm Hg.

[0020] Several forces are developed within the dome and about the rim as a result of evacuating air from the dome. A suction force is developed within the dome 12 equal to the vacuum pressure multiplied by the enclosed tissue surface area 30. The vacuum or vacuum pressure may also be thought of as a negative pressure. The vector sum of the suction force upon the tissue surface area 30 may be called the normal force and is equal to the vacuum pressure multiplied by the normal area 32 of the dome opening, i.e., the area bounded by the periphery 33. An opposing force is imposed on the user by the rim 14 to balance the normal force and is equal but opposite to the normal force. The contact pressure of the rim 14 against the user is equal to this opposing force divided by the annular rim surface area 34, i.e., the surface area between the rim and patient which supports the dome's pressure. Therefore, if the rim surface area 34 is configured to be greater than or equal to the normal area 32 at the dome opening, then the contact pressure against the patient's skin will not exceed the magnitude of the vacuum pressure within the dome 12. Another physical phenomenon further aids in the enlargement forces upon the soft tissue under the dome 12. If the tissue only slightly protrudes into the dome as shown in Figure 3 and as is typically the initial condition, then the surface area 30 under the dome is only slightly larger than the normal area 32 at the dome opening. There-

fore, as the suction force is directly proportional to the surface area of the tissue under the dome, the suction force is only slightly larger than the normal force. As enlargement occurs, more tissue protrudes into the dome 12 as shown in Figure 4 thereby providing more surface area 30 under the dome. Because the surface area 30 under the dome is larger, the suction force generated is increased. Thus, the rate of enlargement increases as treatment continues.

[0021] One specific embodiment includes a dome 12 configured to fit over a human breast as shown in Figures 1 and 2. This embodiment includes a rim 14 having a surface area 34 greater than the normal area 32 of the dome opening thereby preventing medical complications to the soft tissue as long as the pressure is properly regulated within the dome 12. The pressure reducing means 16 is located underneath the patient's breast, so that the apparatus 10 may be hidden under loose-fitting clothes. As with the general embodiment, the vacuum pump assembly 16 of this embodiment is preferably comprised of a vacuum pump 20 with a power source 22, a pressure sensor 24 and servomechanism 26 to drive and control the vacuum pump and to regulate the pressure within the dome 12.

[0022] As shown in Figure 1, this specific embodiment may take the form of a bra 40 having two domes 12 spaced by a hinge 42. Straps 44 may be attached to the bra 40 to retain the bra 40 in place. A gasket 46 may also be included about the rim 14 to improve the patient's comfort and enhance the seal about the rim. In the preferred embodiment, this gasket 46 may be a silicone gel cushion or other soft, conforming type material. Petroleum jelly or other sealant gel may also be used to supplement or supplant the gasket. A manual override 48 is included on the vacuum pump assembly 16 so that the patient or doctor may vary the pressure below the optimal level so as to be more comfortable. Although two vacuum pump assemblies 16 may be used, one depending from each dome 12 so as to provide different pressures in the domes, the preferred embodiment places the domes in fluid communication with a conduit 50.

[0023] A second specific embodiment is shown in Figure 5 wherein the dome 12 is configured to fit over a human penis. As can be seen from the figure, this embodiment comprises essentially the same features as the bra embodiment described above. The principal differences between these embodiments are the configurations of the dome 12' and rim 14' as well as the positioning of the straps 44'.

[0024] As shown in Figure 6, a dome 52 may be conveniently located over an open wound 54. A pump 56 (including an appropriate control) draws a vacuum through a connecting tube 58 in substantially the same manner as has been explained above.

[0025] In order to use the invention, the patient places the dome over the area of desired enlargement and adjusts the straps for comfort. Then the patient simply turns the vacuum pump on and the device goes to work.

These apparatuses are intended to be worn 8-12 hours per day and can be worn during sleep. After several months, notable and long-term enlargement should occur. When the desired enlargement is achieved, the use of the device may be suspended. If additional enlargement is desired, then use may be continued. Occasional use or use at a reduced pressure may also be desired to maintain the desired enlargement.

[0026] For alternate applications, such as in a hospital, clinic or other professional setting, the invention may be applied to the area of desired enlargement, or over an open wound or ulcer, and the vacuum pump and control turned on in order to automatically apply an appropriate regimen of vacuum and rest. As noted above, a vacuum may be developed with the invention and maintained at a continuous negative pressure sufficient to provide tissue enlargement and yet not cause damage to surrounding soft tissue for extended time periods. Alternatively, a "cycling" regimen may be provided by the invention which may promote more rapid tissue enlargement. For example, the vacuum pump may be controlled to develop a pressure as high as 100 mm Hg for several minutes and then return to a much lower level considered to be safe for extended periods, such as between 15-20 or even 35 mm Hg. Upon further testing, other protocols for treatment or use may be found to produce an accelerated enlargement of soft tissue. The present invention should not be considered as limited to any particular protocol as the inventor contemplates that different protocols may be readily learned and utilized with the present invention.

[0027] There are various changes and modifications which may be made to the invention as would be apparent to those skilled in the art. However, these changes or modifications are included in the teaching of the disclosure and it is intended that the invention be limited only by the scope of the claims appended hereto.

Claims

1. An apparatus for promoting enlargement of soft tissue, the apparatus comprising a hollow, fluid-imperious dome (12) having a rigidity that is sufficient to withstand a negative pressure within the dome, pressure reducing means for evacuating air from the dome to provide a suction force and means for regulation of the pressure in the dome, the dome having a periphery (33) surrounding an opening 10 an interior of the dome, the opening to the interior of the dome having a normal area (32), the interior of the dome being dimensioned to cover the soft tissue to be enlarged and enclose said soft tissue within the dome, the dome having a rim (14) that extends around the dome periphery (33) and the rim (14) having a contact surface area (34) for contacting the skin surrounding the soft tissue to be enlarged to support and seal the dome periphery

against the skin, the apparatus **characterized by** the rim contact surface area being equal to or greater than the normal area (32) of the dome opening such that when a negative pressure is drawn inside the dome by operation of the pressure reducing means, a resultant contact pressure exerted by the rim (14) contact surface area against the skin surrounding the soft tissue to be enlarged is less than the negative pressure drawn in the dome thereby minimizing damage to the skin surrounding the soft tissue to be enlarged.

2. An apparatus as claimed in claim 1 **characterized by** a vacuum pump (20) connected in communication with the interior of the dome (12) for reducing pressure within the dome.
3. An apparatus as claimed in claim 2 **characterized by** a regulator (24, 26) connected to the vacuum pump (20) for maintaining a desired negative pressure within the dome.
4. An apparatus as claimed in any one of claims 1 to 3 **characterized by** a gasket (46) provided on the dome rim (14) for providing comfort and sealing of the dome (12).
5. An apparatus as claimed in claim 2, 3 or 4, **characterized by** a manual control (48) on the vacuum pump (20) provided for user adjustment of the pressure within the dome.
6. An apparatus as claimed in any one of the preceding claims **characterized by** the dome (12) being one of a pair of domes (12), each dome having a rim (14) with a contact surface area (34); a hinge (42) joining the rims of the pair of domes; and a fluid communication means (5) connected between the pair of domes to equalize vacuum pressure within the pair of domes.
7. An apparatus as claimed in claim 6 **characterized by** each dome (12) of the pair of domes being shaped to enclose a human breast in the dome interior.
8. An apparatus as claimed in claims 3, 4 or 5 **characterized by** the dome (12) being one of a pair of domes (12), each dome having a rim (14) within a contact surface area (34); a hinge (43) joining the rims of the pair of domes; the vacuum pump (20) being one of a pair of vacuum pumps (20), each pump being connected in communication with the interior of one of the pair of domes; and the regulator (24, 26) being one of a pair of regulators (24, 26), each regulator being connected to one of the pair of pumps (20) for maintaining a desired vacuum pressure in each dome.

9. An apparatus as claimed in of any of claims 1 to 5 **characterized by** the dome (12) being shaped to enclose a human penis in the dome interior.
10. An apparatus as claimed in of any one of claims 1 to 5 **characterized by** the soft tissue to be enlarged has a wound (54) and the dome (12) is shaped to enclose the wound in the interior of the dome.
11. An apparatus as claimed in any one of the preceding claims **characterized by** the apparatus is manually portable.
12. An apparatus as claimed in any one of the preceding claims **characterized by** the rim contact surface area (34) being dimensioned sufficiently large to limit the contact pressure of the rim with the soft tissue that surrounds the soft tissue that is to be enlarged to less than 20 mm Hg when the negative pressure is created in the dome.
13. An apparatus as claimed in claim 12 **characterized by** the negative pressure being between 25 mm of Hg and 35 mm of Hg.

Patentansprüche

1. Vorrichtung zur Förderung der Weichgewebevergrößerung, wobei die Vorrichtung eine hohle, flüssigkeitsdichte Kuppel (12) mit einer ausreichenden Steifigkeit, um einem Unterdruck innerhalb der Kuppel widerstehen zu können. Druckminderungsmittel zum Evakuieren von Luft aus der Kuppel, um eine Sogkraft in der Kuppel bereit zu stellen, und Mittel zur Regulierung des Drucks innerhalb der Kuppel aufweist, wobei die Kuppel einen Umfang (33) aufweist, der eine Öffnung in das Innere der Kuppel umgibt, und die Öffnung in das Innere der Kuppel einen Normalbereich (32) aufweist, wobei das Innere der Kuppel so dimensioniert ist, dass es das zu vergrößernde Weichgewebe überdeckt und das Weichgewebe innerhalb der Kuppel umgibt, wobei die Kuppel einen Rand (14) aufweist, der sich um den Umfang (33) der Kuppel herum erstreckt, und wobei der Rand (14) einen Auflageflächenbereich (34) zum Kontaktieren der das zu vergrößernde Weichgewebe umgebenden Haut aufweist, um den Kuppelumfang gegen die Haut abzustützen und abzudichten, wobei die Vorrichtung **dadurch gekennzeichnet ist, dass** der Randauflageflächenbereich gleich oder größer als der Normalbereich (32) der Kuppelöffnung ist, so dass, wenn durch Betätigung der Druckminderungsmittel ein Unterdruck in das Innere der Kuppel gesaugt wird, ein durch den Rand-(14)-auflageflächenbereich gegen die das zu vergrößernde Weichgewebe umschließende Haut ausgeübter resultierender An-

pressdruck kleiner ist als der in die Kuppel gesaugte Unterdruck, wodurch eine Schädigung der das zu vergrößernde Weichgewebe umschließenden Haut minimiert wird.

2. Vorrichtung nach Anspruch 1, **gekennzeichnet durch** eine mit dem Inneren der Kuppel (12) in Verbindung stehende Vakuumpumpe (20) zur Verminderung des Drucks innerhalb der Kuppel.
3. Vorrichtung nach Anspruch 2, **gekennzeichnet durch** ein mit der Vakuumpumpe (20) verbundenes Regelgerät (24, 26) zur Aufrechterhaltung eines gewünschten Unterdrucks innerhalb der Kuppel.
4. Vorrichtung nach einem der Ansprüche 1 bis 3, **gekennzeichnet durch** einen auf dem Kuppelrand (14) angeordneten Dichttring (46) zum Bereitstellen von Komfort und zum Abdichten der Kuppel (12).
5. Vorrichtung nach einem der Ansprüche 2, 3 oder 4, **gekennzeichnet durch** eine an der Vakuumpumpe (20) zur Einstellung des Druckes innerhalb der Kuppel **durch** den Benutzer vorgesehene Handbedienung (48).
6. Vorrichtung nach einem der vorstehenden Ansprüche, **gekennzeichnet dadurch, dass** die Kuppel (12) eine eines Paares von Kuppeln (12) darstellt, wobei jede Kuppel einen Rand (14) mit einem Kontaktflächenbereich (34) aufweist: durch ein die Ränder des Paares von Kuppeln verbindendes Band (42); und zwischen dem Paar von Kuppeln angeschlossene Flüssigkeitsverbindungsmittel (50), um den Unterdruck innerhalb des Paares von Kuppeln anzugleichen.
7. Vorrichtung nach Anspruch 6, **dadurch gekennzeichnet, dass** jede Kuppel (12) des Paares von Kuppeln so geformt ist, dass sie eine menschliche Brust im Kuppelinneren umgeben kann.
8. Vorrichtung nach Anspruch 3, 4 oder 5, **gekennzeichnet dadurch, dass** die Kuppel (12) eine eines Paares von Kuppeln (12) darstellt, wobei jede Kuppel einen Rand (14) innerhalb eines Kontaktflächenbereiches (34) aufweist; durch ein die Ränder des Paares von Kuppeln verbindendes Band (42); dadurch, dass die Vakuumpumpe (20) eine eines Paares von Vakuumpumpen (20) darstellt, wobei jede Pumpe mit dem Inneren einer des Paares von Kuppeln in Verbindung steht; und dadurch, dass das Regelgerät (24, 26) eines eines Paares von Regelgeräten (24, 26) darstellt, wobei jedes Regelgerät mit einer des Paares von Pumpen (20) verbunden ist, um einen gewünschten Unterdruck in jeder Kuppel aufrecht zu erhalten.

9. Vorrichtung nach einem der Ansprüche 1 bis 5, **dadurch gekennzeichnet, dass** die Kuppel (12) so geformt ist, dass sie einen menschlichen Penis im Kuppelinneren umgeben kann.
10. Vorrichtung nach einem der Ansprüche 1 bis 5, **dadurch gekennzeichnet, dass** das zu vergrößern- de Weichgewebe eine Wunde (54) aufweist, und dass die Kuppel (12) so geformt ist, dass sie die Wunde im Inneren der Kuppel umgeben kann.
11. Vorrichtung nach einem der vorstehenden Ansprüche, **dadurch gekennzeichnet, dass** die Vorrichtung von Hand tragbar ist.
12. Vorrichtung nach einem der vorstehenden Ansprüche, **dadurch gekennzeichnet, dass** der Rand- kontaktflächenbereich (34) ausreichend groß dimensioniert ist, um den Anpreßdruck des Randes an das Weichgewebe, dass das zu vergrößern- de Weichgewebe umgibt, auf weniger als 20 mm Quecksilbersäule zu begrenzen, wenn der Unter- druck in der Kuppel erzeugt wird.
13. Vorrichtung nach Anspruch 12, **dadurch gekenn- zeichnet, dass** der Unterdruck zwischen 25 mm und 35 mm Quecksilbersäule liegt.

Revendications

1. Appareil pour favoriser l'augmentation de tissus mous, l'appareil comprenant un dôme creux (12) imperméable aux fluides ayant une rigidité qui est suffisante pour supporter une pression négative à l'intérieur du dôme, un moyen de réduction de pression pour évacuer l'air du dôme afin de fournir une force de succion et un moyen pour la régulation de la pression dans le dôme, le dôme ayant une périphérie (33) entourant une ouverture menant à l'intérieur du dôme, l'ouverture menant à l'intérieur du dôme ayant une aire normale (32), l'intérieur du dôme étant dimensionné de façon à couvrir le tissu mou à augmenter et à enfermer ledit tissu mou à l'intérieur du dôme, le dôme ayant un rebord (14) qui s'étend autour de la périphérie (33) du dôme et le rebord (14) ayant une aire de surface de contact (34) pour le contact avec la peau entourant le tissu mou à augmenter afin de supporter et de fermer hermétiquement la périphérie du dôme contre la peau, l'appareil étant **caractérisé par le fait que** l'aire de surface de contact du rebord est supérieure ou égale à l'aire normale (32) de l'ouverture du dôme de telle sorte que lorsqu'une pression négative est créée à l'intérieur du dôme par le biais du moyen de réduction de pression, une pression de contact résultante exercée par l'aire de surface de contact du rebord (14) sur la peau entourant le tissu mou à

augmenter est inférieure à la pression négative créée dans le dôme, minimisant de ce fait les dégâts causés à la peau qui entoure le tissu mou à augmenter.

2. Appareil selon la revendication 1, **caractérisé par** une pompe à vide (20) connectée en communication avec l'intérieur du dôme (12) pour réduire la pression à l'intérieur du dôme.
3. Appareil selon la revendication 2, **caractérisé par** un régulateur (24, 26) connecté à la pompe à vide (20) pour maintenir une pression négative souhaitée à l'intérieur du dôme.
4. Appareil selon l'une quelconque des revendications 1 à 3, **caractérisé par** un joint (46) placé sur le rebord (14) du dôme pour assurer le confort et l'étanchéité du dôme (12).
5. Appareil selon la revendication 2, 3 ou 4, **caracté- risé par** une commande manuelle (48) sur la pompe à vide (20) pour permettre le réglage par l'utili- sateur de la pression régnant à l'intérieur du dôme.
6. Appareil selon l'une quelconque des revendications précédentes, **caractérisé par le fait que** le dôme (12) est un dôme d'une paire de dômes (12), cha- que dôme ayant un rebord (14) muni d'une aire de surface de contact (34), une charnière (42) reliant les rebords de la paire de dômes, et un moyen de communication (5) de fluide connecté entre les dô- mes de la paire pour égaliser la pression de vide dans la paire de dômes.
7. Appareil selon la revendication 6, **caractérisé par le fait que** chaque dôme (12) de la paire de dômes a une forme prévue pour enfermer un sein humain à l'intérieur du dôme.
8. Appareil selon la revendication 3, 4 ou 5, **caracté- risé par le fait que** le dôme (12) est un dôme d'une paire de dômes (12), chaque dôme ayant un rebord (14) muni d'une aire de surface de contact (34), une charnière (43) reliant les rebords de la paire de dô- mes, la pompe à vide (20) étant une pompe d'une paire de pompes à vide (20), chaque pompe étant connectée en communication avec l'intérieur de l'un des dômes de la paire de dômes, et le régulateur (24, 26) étant un régulateur d'une paire de régula- teurs (24, 26), chaque régulateur étant connecté à l'une des pompes de la paire de pompes (20) pour maintenir une pression de vide souhaitée dans cha- que dôme.
9. Appareil selon l'une quelconque des revendications 1 à 5, **caractérisé par le fait que** le dôme (12) a une forme prévue pour enfermer un pénis humain

à l'intérieur du dôme.

10. Appareil selon l'une quelconque des revendications 1 à 5, **caractérisé par le fait que** le tissu mou à augmenter comporte une plaie (54) et que le dôme (12) a une forme prévue pour enfermer la plaie à l'intérieur du dôme. 5
11. Appareil selon l'une quelconque des revendications précédentes, **caractérisé par le fait que** l'appareil est portatif manuellement. 10
12. Appareil selon l'une quelconque des revendications précédentes, **caractérisé par le fait que** l'aire de surface de contact (34) est dimensionnée de façon à être suffisamment grande pour limiter la pression de contact du rebord avec le tissu mou qui entoure le tissu mou à augmenter à moins de 20 mm Hg quand la pression négative est créée à l'intérieur du dôme. 15 20
13. Appareil selon la revendication 12, **caractérisé par le fait que** la pression négative est comprise entre 25 mm de Hg et 35 mm de Hg. 25

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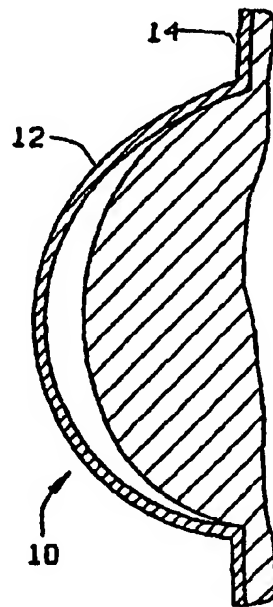
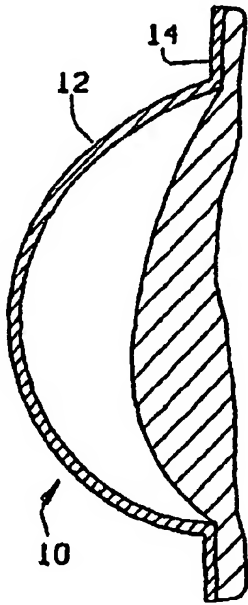
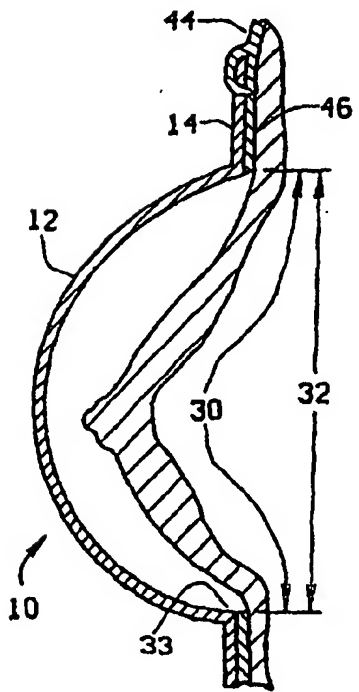
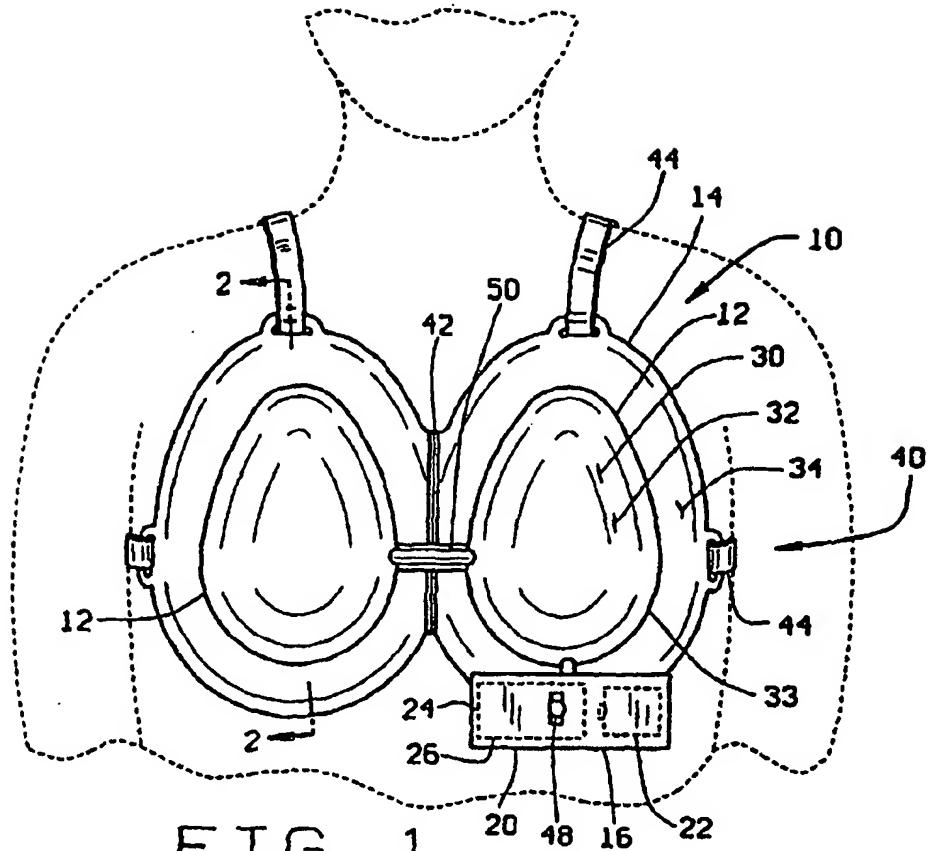
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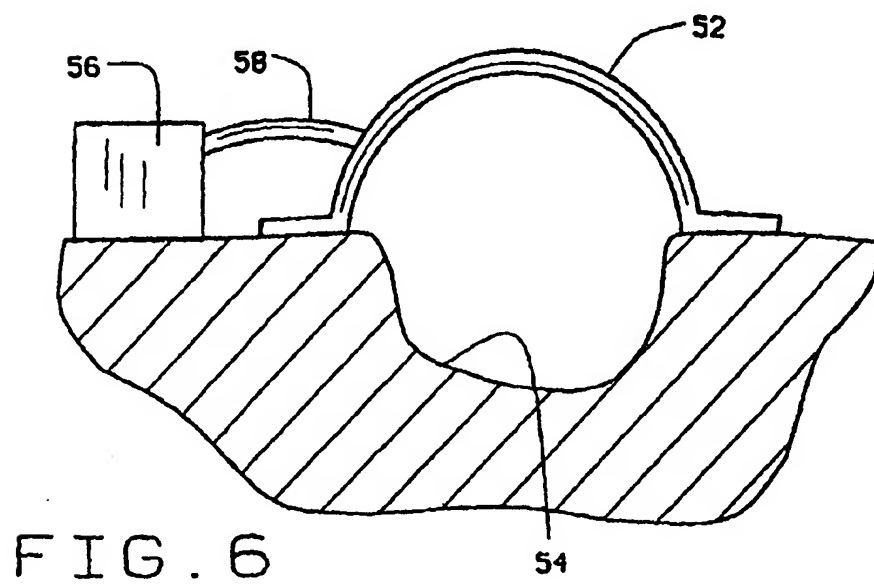
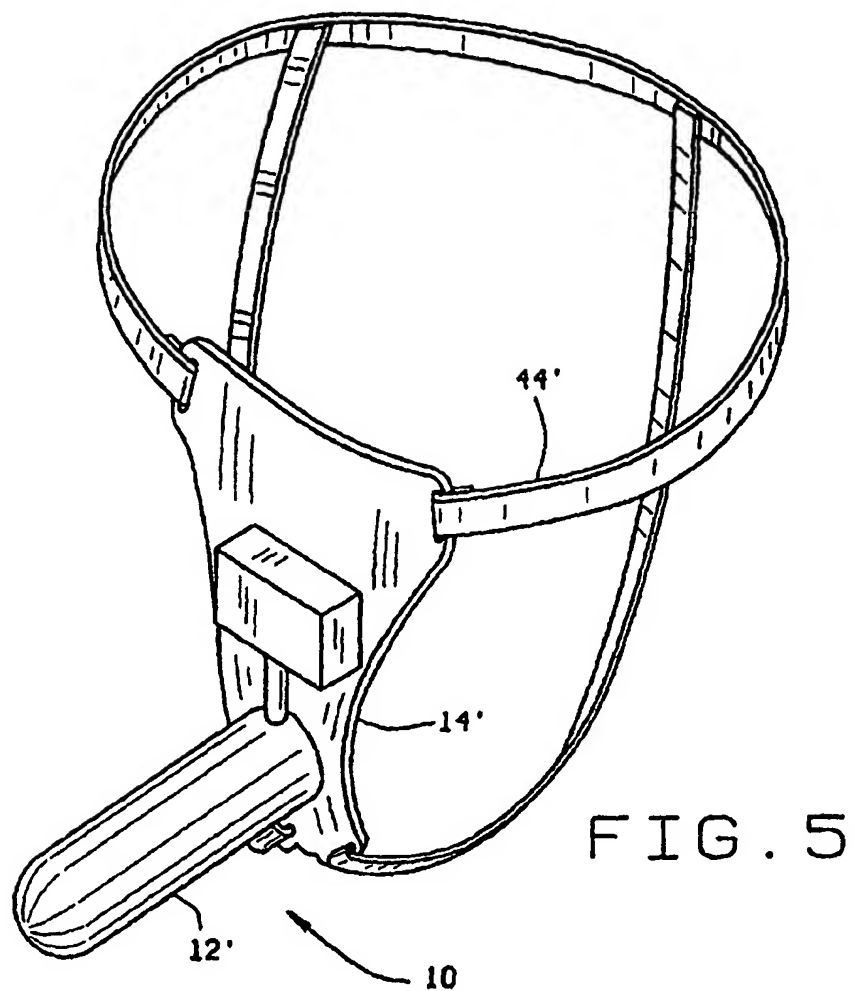
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